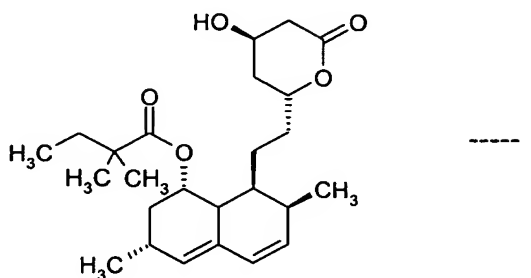


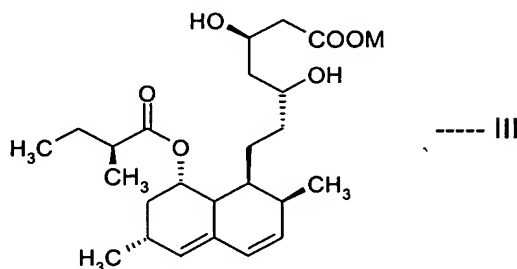
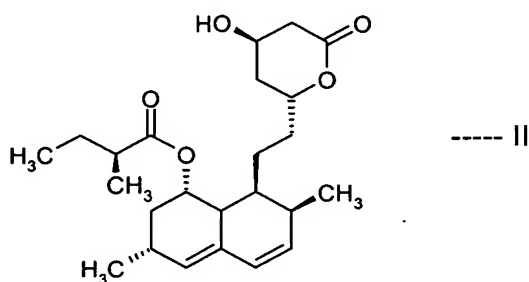
We claim:

1) A process for the preparation of simvastatin of formula I:



which comprises the steps of:

5 a) reacting compound of formula II (lovastatin) or formula III:



wherein M is H, metal ion or NH<sub>4</sub>,

with the compound of formula IV:

10 HNR<sub>1</sub>R<sub>2</sub> ----- IV

wherein

R<sub>1</sub> is -R<sub>5</sub>-X-R<sub>6</sub> wherein

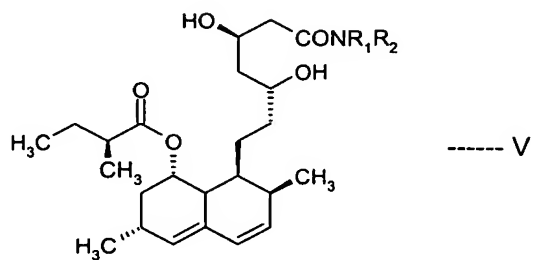
R<sub>5</sub> is alkyl, arylalkyl or cycloalkyl,

X is O or S and

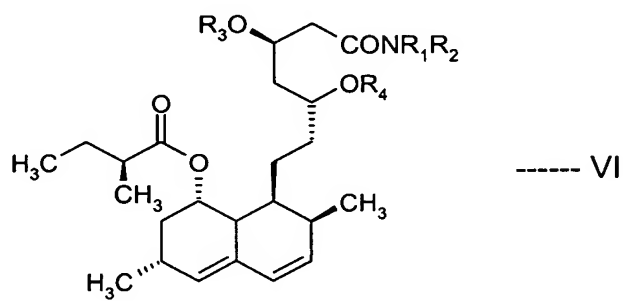
15 R<sub>6</sub> is alkyl, arylalkyl, cycloalkyl or aryl; and

R<sub>2</sub> is independently selected from H, alkyl, cycloalkyl, arylalkyl and a group as defined for R<sub>1</sub>;

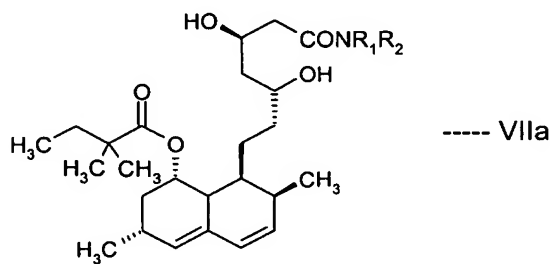
or R<sub>1</sub> and R<sub>2</sub> may be bonded to form a cyclic ether or cyclic thio ether;  
to produce a compound of formula V:

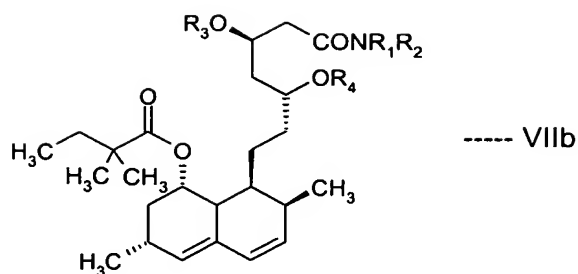


- 5 wherein R<sub>1</sub> and R<sub>2</sub> are as defined above,  
(b) optionally protecting the two hydroxyl groups of the said compound of the formula V  
to produce a compound of the formula VI:



- wherein R<sub>3</sub> and R<sub>4</sub> represents suitable protecting groups,  
10 (c) methylating the said compound of formula V or VI to give a compound of formula  
VIIa or VIIb:

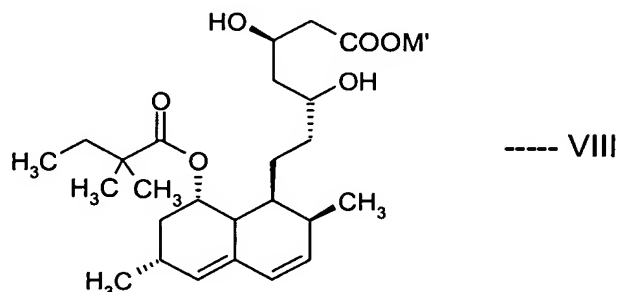




wherein  $R_1$ ,  $R_2$ ,  $R_3$  and  $R_4$  are as defined above,

- 5 (d) hydrolyzing the amide group if the product of the above step is the said compound of formula VIIa or deprotecting the two protected hydroxy groups prior to hydrolysis if the product of the above step is the said compound of formula VIIb, optionally treating the hydrolyzed product with aqueous ammonia, to produce a compound of formula VIII:

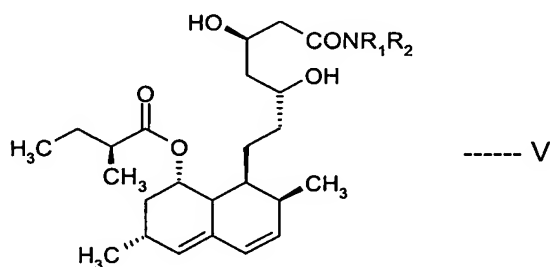
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wherein  $M'$  is a metal such as sodium or potassium or  $NH_4$ ,

- (e) lactonizing the said compound of formula VIII to produce simvastatin of formula I.
- 2) A process according to claim 1, wherein the hydroxy groups are not protected before methylation.
- 15 3) A process according to claim 1 and 2, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl, and  $R_2$  is selected from H, methoxyethyl, ethoxyethyl and methoxymethyl.
- 4) A process according to claim 3, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl, and  $R_2$  is H.
- 20 5) A process according to claim 1 - 4, wherein  $R_1$  is methoxyethyl and  $R_2$  is H.
- 6) A process according to claim 1, wherein methylation is carried out using an alkali metal amide and a methyl halide.

- 7) A process according to claim 6, wherein the alkali metal is lithium, sodium or potassium; and the methyl halide is methyl iodide, methyl chloride or methyl bromide.
- 8) A process according to claim 6 and 7, wherein the alkali metal amide is lithium pyrrolidide and the methylhalide is methyl iodide.
- 9) A process according to claim 1, wherein the starting compound is lovastatin of formula II.
- 10) A process according to claim 1, wherein  $R_3$  and  $R_4$  represent silyl protecting groups.
- 11) A process according to claim 10, wherein the silyl protecting groups are selected from t-butyldimethylsilyl and trimethylsilyl groups.
- 12) A process according to claim 1, wherein i) lovastatin is treated with methoxyethyl amine in an organic solvent to produce the compound of the formula V wherein  $R_1$  is methoxyethyl- and  $R_2$  is H, ii) methylating the product obtained in the previous step with lithium pyrrolidide in tetrahydrofuran and methyl iodide to produce the compound of the formula VIIa wherein  $R_1$  is methoxyethyl- and  $R_2$  is H, iii) hydrolyzing the product obtained in the previous step with a strong base to obtain the compound of the formula VIII, iv) adding aqueous ammonia to the product obtained in the previous step to produce simvastatin ammonium salt, and v) lactonizing the product obtained in the previous step to produce simvastatin.
- 13) A compound of the formula V:



wherein

$R_1$  is  $-R_5-X-R_6$  wherein

$R_5$  is alkyl, arylalkyl or cycloalkyl,

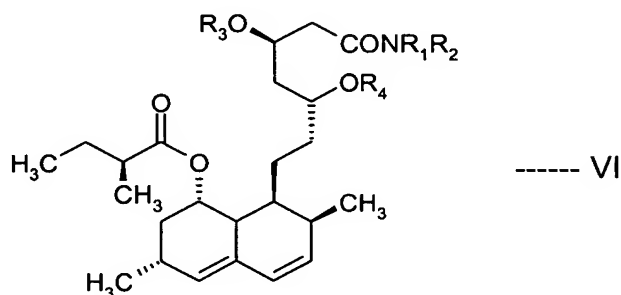
$X$  is O or S and

$R_6$  is alkyl, arylalkyl, cycloalkyl or aryl; and

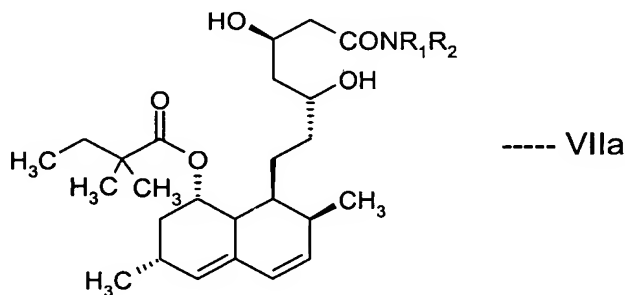
$R_2$  is independently selected from H, alkyl, cycloalkyl, arylalkyl and a group as defined for  $R_1$ ;

or  $R_1$  and  $R_2$  may be bonded to form a cyclic ether or cyclic thio ether;

- 14) The compound of the claim 13, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl, and  $R_2$  is selected from H, methoxyethyl, ethoxyethyl and methoxymethyl.
- 15) The compound of claim 14, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl, and  $R_2$  is H.
- 16) The compound of claim 15, wherein  $R_1$  is methoxyethyl and  $R_2$  is H.
- 17) A compound of the formula VI:



- 18) wherein  $R_1$  and  $R_2$  are as defined in formula V of claim 13; and  $R_3$  and  $R_4$  represents suitable protecting groups.
- 19) The compound of claim 17, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl,  $R_2$  is selected from H, methoxyethyl, ethoxyethyl and methoxymethyl and  $R_3$  and  $R_4$  are selected from silyl protecting groups such as t-butyltrimethylsilyl and trimethylsilyl groups.
- 20) The compound of claim 17 or 18, wherein  $R_1$  is selected from methoxyethyl, ethoxyethyl and methoxymethyl, and  $R_2$  is H.
- 21) The compound of claim 19, wherein  $R_1$  is methoxyethyl and  $R_2$  is H.
- 22) The compound of the formula VIIa:



wherein  $R_1$  and  $R_2$  are as defined in the formula V of claim 13.

